

Prisca 5.1.0.17
Date of report: 21-02-2025

NA

Patient data						
Name	W/O RANK /NK AJEET KUMAR		Patient ID	0442502180071		
Birthday	10-12-1998		Sample ID	A0611468		
Age at sample date	26.2		Sample Date	18-02-2025		
Gestational age	13 + 2					
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	50	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age	13 + 2		
PAPP-A	4.4 mIU/mL	0.67	Method	CRL Robinson		
fb-hCG	27.36 ng/mL	0.75	Scan date	18-02-2025		
Risks at sampling date			Crown rump length in mm		74	
Age risk	1:928		Nuchal translucency MoM	0.56		
Biochemical T21 risk	1:4395		Nasal bone	present		
Combined trisomy 21 risk	<1:10000		Sonographer	NA		
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD		
Risk			Trisomy 21			
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!			
Trisomy 13/18 + NT						
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.						

Sign of Physician